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What Is Claimed Is:

1. A compound of Formula I:

wherein each of R^1 and R^{11} is a group independently selected from hydrido, alkyl, alkylaminoalkyl and phenyl; wherein p is a number selected from zero through five, inclusive; wherein r is a number selected from zero, one and two; wherein R^2 is selected from hydrido and alkyl; wherein R^3 is a group selected from hydrido, cycloalkylalkyl, aralkyl and haloaralkyl; wherein each of R^4 and R^6 is a group independently selected from hydrido and methyl; wherein R^5 is selected from

wherein V is selected from hydrido, alkyl, cycloalkyl, aryl and aralkyl; wherein each of R^9 and \tilde{R}^{10} is a group independently selected from hydrido, alkyl, alkenyl, alkynyl, cycloalkyl and aryl; wherein m is a number selected from zero through three; wherein n is a number selected from zero through three; wherein R^7 is a group selected from alkyl, cycloalkylalkyl and aralkyl; wherein R^8 is a group selected from hydrido, alkyl, hydroxyalkyl, cycloalkyl, cycloalkylalkyl, alkenyl and haloalkenyl; wherein each of R^{12} and R^{13} is a group independently selected from hydrido, alkyl, cycloalkyl,

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cycloalkylalkyl, alkylacyl, aryl, aralkyl, haloaryl and haloaralkyl; and wherein any one of said R¹ through R¹³ groups having a substitutable position may be substituted with one or more groups selected from alkyl, hydroxy, hydroxyalkyl, halo, alkoxy, alkoxyalkyl and alkenyl; or a pharmaceutically-acceptable salt thereof.

2. Compound of Claim 1 wherein each of R^1 and R^{11} is independently selected from hydrido, methyl, ethyl, n-propyl, isopropyl, n-butyl, sec-butyl, iso-butyl, tert-butyl, N,N'-dimethylaminomethyl, N,N'-diethylaminomethyl, N,N'diethylaminoethyl and phenyl; wherein p is a number selected from zero through four, inclusive; wherein r is a number selected from zero, one and two; wherein R^2 is selected from hydrido and alkyl; wherein R^3 is selected from hydrido, cycloalkylalkyl, phenylalkyl, halophenylalkyl, naphthylalkyl and halonaphthylalkyl; wherein each of R^4 and R^6 is independently selected from hydrido and methyl; wherein R^5 is selected from

$$\frac{\left(CH_{2} \right)^{-}}{m} \frac{\left(CH_{2} \right)^{-}}{\left(CH_{2} \right)^{-}} = C \equiv C-V$$

wherein V is selected from hydrido, alkyl, phenyl and benzyl; wherein each of R⁹ and R¹⁰ is independently selected from hydrido, alkyl, alkenyl, alkynyl, cycloalkyl and aryl; wherein m is a number selected from zero through three; wherein n is a number selected from zero through three; wherein R⁷ is selected from cyclohexylmethyl and benzyl, either one of which may be substituted with one or more groups selected from alkyl, hydroxy and alkoxy; wherein R⁸ is selected from hydrido, alkyl, cycloalkyl, cycloalkylalkyl, hydroxyalkyl, alkenyl

and haloalkenyl; and wherein each of R¹² and R¹³ is independently selected from hydrido, alkyl, cycloalkyl, cycloalkyl, alkanoyl, halophenyl, phenylalkyl, halophenylalkyl, naphthyl, halonaphthyl, naphthylalkyl and halonaphthylalkyl; or a pharmaceutically-acceptable salt thereof.

Compound of Claim 2 wherein each of R1 and R11 is independently selected from hydrido, methyl, ethyl, n-propyl and isopropyl; wherein p is a number selected 10 from zero through three, inclusive; wherein r is a number selected from zero, one and two; wherein R^2 is selected from hydrido, methyl, ethyl and n-propyl; wherein R^3 is selected from hydrido, cyclohexylmethyl, benzyl, phenylethyl, fluorobenzyl, fluorophenylethyl, 15 chlorobenzyl, chlorophenylethyl, naphthylmethyl, naphthylethyl, fluoronaphthylmethyl and chloronaphthylmethyl; wherein each of R^4 and R^6 is independently selected from hydrido and methyl; wherein R⁵ is selected from 20

$$-(CH2)m C - C = C-V$$

wherein V is selected from hydrido, alkyl, cycloalkyl, aryl and aralkyl; wherein m is a number selected from one through three; wherein R⁷ is cyclohexylmethyl; wherein R⁸ is selected from methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, sec-butyl, tert-butyl, cyclopropyl, cyclobutyl, cyclopropylmethyl, cyclobutylmethyl, cyclohexylmethyl, allyl and vinyl; and wherein each of R¹² and R¹³ is independently selected from hydrido, methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, sec-butyl,

tert-butyl, cyclopropyl, cyclopropylmethyl,
cyclopropylethyl, propylcarbonyl, ethylcarbonyl,
methylcarbonyl, phenyl, benzyl, phenylethyl,
monochlorophenyl, dichlorophenyl, monofluorophenyl,
difluorophenyl, monochlorophenylmethyl,
monochlorophenylethyl, dichlorophenylmethyl,
dichlorophenylethyl, naphthyl, monofluoronaphthyl,
monochloronaphthyl, naphthylmethyl, naphthylethyl,
fluoronapthylmethyl and chloronaphthylethyl; or a
pharmaceutically-acceptable salt thereof.

4. Compound of Claim 3 wherein each of R¹ and R¹¹ is independently hydrido or methyl; wherein p is a number selected from zero through three, inclusive; wherein r is zero or two; wherein R² is selected from hydrido, methyl, ethyl and n-propyl; wherein R³ is selected from hydrido, cyclohexylmethyl, benzyl, phenylethyl, phenylpropyl, fluorobenzyl, fluorophenylethyl, chlorobenzyl, chlorophenylethyl, naphthylmethyl, fluoronaphthylmethyl and chloronaphthylmethyl; wherein each of R⁴ and R⁶ is hydrido; wherein R⁵ is selected from

$$--(CH_2)_m$$
 $C \equiv C-V$

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wherein V is selected from hydrido and methyl; wherein m is one or two; wherein R⁷ is cyclohexylmethyl; wherein R⁸ is selected from ethyl, n-propyl, n-butyl, isobutyl, cyclopropyl, cyclobutyl, cyclopropylmethyl, allyl and vinyl; wherein each of R¹² and R¹³ is independently selected from hydrido, methyl, ethyl, n-propyl, isopropyl, cyclopropylmethyl, phenyl, benzyl, monochlorophenyl and dichlorophenyl; or a pharmaceutically-acceptable salt thereof.

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5. Compound of Claim 4 of Formula II

wherein r is zero or two; wherein q is two or three; wherein R² is selected from hydrido, methyl, ethyl and phenyl; wherein R³ is selected from hydrido, cyclohexylmethyl, benzyl, fluorobenzyl, chlorobenzyl, fluoronaphthylmethyl and chloronaphthylmethyl; wherein each of R⁴ and R⁶ is hydrido; wherein R⁵ is selected from

$$--(CH_2)_m$$
 $C \equiv C-V$

wherein V is selected from hydrido and methyl; wherein m is one or two; wherein R⁷ is cyclohexylmethyl; wherein R⁸ is selected from n-propyl, isobutyl, cyclopropyl, cyclopropylmethyl, allyl and vinyl; wherein R¹² and R¹³ is independently selected from methyl, ethyl and isopropyl; or a pharmaceutically-acceptable salt thereof.

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6. Compound of Claim 5 selected from compounds, their tautomers and pharmaceutically-acceptable salts thereof, of the group consisting of:

7. Compound of Claim 6 which is N-[1R*-[[[1S,1R*-(cyclohexylmethyl)-2S*,3R*-dihydroxy-5-

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methylhexyl]amino]carbonyl]-3-butynyl]-αR*-[[[2-(dimethylamino)ethyl]sulfonyl]methyl]benzenepropanamide or a pharmaceutically-acceptalble salt thereof.

8. A pharmaceutical composition comprising a therapeutically-effective amount of a renin-inhibiting compound and a pharmaceutically-acceptable carrier or diluent, said renin-inhibiting compound selected from a family of compounds of Formula I:

wherein each of R^1 and R^{11} is a group independently selected from hydrido, alkyl, alkylaminoalkyl and phenyl; wherein p is a number selected from zero through five, inclusive; wherein r is a number selected from zero, one and two; wherein R^2 is selected from hydrido and alkyl; wherein R^3 is a group selected from hydrido, cycloalkylalkyl, aralkyl and haloaralkyl; wherein each of R^4 and R^6 is a group independently selected from hydrido and methyl; wherein R^5 is selected from

$$-(CH2) - C = C-V$$

$$R^{10}$$

wherein V is selected from hydrido, alkyl, cycloalkyl, aryl and aralkyl; wherein each of R⁹ and R¹⁰ is a group independently selected from hydrido, alkyl, alkenyl, alkynyl, cycloalkyl and aryl; wherein m is a number selected from zero through three; wherein n is a number

selected from zero through three; wherein R⁷ is a group selected from alkyl, cycloalkylalkyl and aralkyl; wherein R⁸ is a group selected from hydrido, alkyl, hydroxyalkyl, cycloalkyl, cycloalkylalkyl, alkenyl and haloalkenyl; wherein each of R¹² and R¹³ is a group independently selected from hydrido, alkyl, cycloalkyl, cycloalkylalkyl, alkylacyl, aryl, aralkyl, haloaryl and haloaralkyl; and wherein any one of said R¹ through R¹³ groups having a substitutable position may be substituted with one or more groups selected from alkyl, hydroxy, hydroxyalkyl, halo, alkoxy, alkoxyalkyl and alkenyl; or a pharmaceutically-acceptable salt thereof.

The composition of Claim 8 wherein each of 9. ${\bf R}^1$ and ${\bf R}^{11}$ is independently selected from hydrido, methyl, 15 ethyl, n-propyl, isopropyl, n-butyl, sec-butyl, isobutyl, tert-butyl, N,N'-dimethylaminomethyl, N,N'diethylaminomethyl, N,N'diethylaminoethyl and phenyl; wherein p is a number selected from zero through four, inclusive; wherein r is a number selected from zero, one 20 and two; wherein R² is selected from hydrido and alkyl; wherein R³ is selected from hydrido, cycloalkylalkyl, phenylalkyl, halophenylalkyl, naphthylalkyl and halonaphthylalkyl; wherein each of R^4 and R^6 is independently selected from hydrido and methyl; wherein 25 R^5 is selected from

$$--(CH2) - C = C-V$$

$$R^{10}$$

wherein V is selected from hydrido, alkyl, phenyl and benzyl; wherein each of R^9 and R^{10} is independently selected from hydrido, alkyl, alkenyl, alkynyl,

cycloalkyl and aryl; wherein m is a number selected from zero through three; wherein n is a number selected from zero through three; wherein R⁷ is selected from cyclohexylmethyl and benzyl, either one of which may be substituted with one or more groups selected from alkyl, hydroxy and alkoxy; wherein R⁸ is selected from hydrido, alkyl, cycloalkyl, cycloalkylalkyl, hydroxyalkyl, alkenyl and haloalkenyl; and wherein each of R¹² and R¹³ is independently selected from hydrido, alkyl, cycloalkyl, cycloalkyl, cycloalkyl, halophenyl, phenylalkyl, halophenylalkyl, naphthyl, halonaphthyl, naphthylalkyl and halonaphthylalkyl; or a pharmaceutically-acceptable salt thereof.

10. The composition of Claim 9 wherein each of R¹ and R¹¹ is independently selected from hydrido, methyl, ethyl, n-propyl and isopropyl; wherein p is a number selected from zero through three, inclusive; wherein r is a number selected from zero, one and two; wherein R² is selected from hydrido, methyl, ethyl and n-propyl; wherein R³ is selected from hydrido, cyclohexylmethyl, benzyl, phenylethyl, fluorobenzyl, fluorophenylethyl, chlorobenzyl, chlorophenylethyl, naphthylmethyl, naphthylethyl, fluoronaphthylmethyl and chloronaphthylmethyl; wherein each of R⁴ and R⁶ is independently selected from hydrido and methyl; wherein R⁵ is selected from

$$-(CH2) = \begin{bmatrix} R9 \\ C \\ R10 \end{bmatrix} - C \equiv C-V$$

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wherein V is selected from hydrido, alkyl, cycloalkyl, aryl and aralkyl; wherein m is a number selected from one

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through three; wherein R⁷ is cyclohexylmethyl; wherein R⁸ is selected from methyl, ethyl, n-propyl, isopropyl, nbutyl, isobutyl, sec-butyl, tert-butyl, cyclopropyl, cyclobutyl, cyclopropylmethyl, cyclobutylmethyl, cyclohexylmethyl, allyl and vinyl; and wherein each of R12 and R13 is independently selected from hydrido, methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, sec-butyl, tert-butyl, cyclopropyl, cyclopropylmethyl, cyclopropylethyl, propylcarbonyl, ethylcarbonyl, methylcarbonyl, phenyl, benzyl, phenylethyl, 10 monochlorophenyl, dichlorophenyl, monofluorophenyl, difluorophenyl, monochlorophenylmethyl, monochlorophenylethyl, dichlorophenylmethyl, dichlorophenylethyl, naphthyl, monofluoronaphthyl, monochloronaphthyl, naphthylmethyl, naphthylethyl, 15 fluoronapthylmethyl and chloronaphthylethyl; or a pharmaceutically-acceptable salt thereof.

of R¹ and R¹¹ is independently hydrido or methyl; wherein p is a number selected from zero through three, inclusive; wherein r is zero or two; wherein R² is selected from hydrido, methyl, ethyl and n-propyl; wherein R³ is selected from hydrido, cyclohexylmethyl, benzyl, phenylethyl, phenylpropyl, fluorobenzyl, fluorophenylethyl, chlorobenzyl, chlorophenylethyl, naphthylmethyl, fluoronaphthylmethyl and chloronaphthylmethyl; wherein each of R⁴ and R⁶ is hydrido; wherein R⁵ is selected from

 $--(CH_2)_m$ $C \equiv C-V$

wherein V is selected from hydrido and methyl; wherein m is one or two; wherein R^7 is cyclohexylmethyl; wherein R^8 is selected from ethyl, n-propyl, n-butyl, isobutyl, cyclopropyl, cyclobutyl, cyclopropylmethyl, allyl and vinyl; wherein each of R^{12} and R^{13} is independently

selected from hydrido, methyl, ethyl, n-propyl, isopropyl, cyclopropylmethyl, phenyl, benzyl, monochlorophenyl and dichlorophenyl; or a pharmaceutically-acceptable salt thereof.

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The composition of Claim 11 wherein said compound is of Formula II

$$R^{13} = \begin{bmatrix} R^{13} & 0 & R^{5} & H & OH \\ N & N - [CH_2]_q & S[O]_r & R^{3} & H & OH \\ R^{12} & OH & OH \end{bmatrix}$$

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wherein r is zero or two; wherein q is two or three; wherein R^2 is selected from hydrido, methyl, ethyl and phenyl; wherein R³ is selected from hydrido, cyclohexylmethyl, benzyl, fluorobenzyl, chlorobenzyl, fluoronaphthylmethyl and chloronaphthylmethyl; wherein each of R^4 and R^6 is hydrido; wherein R^5 is selected from

$$--(CH_2)_m$$
 $C \equiv C-V$

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wherein V is selected from hydrido and methyl; wherein m is one or two; wherein R^7 is cyclohexylmethyl; wherein R^8 is

selected from n-propyl, isobutyl, cyclopropyl, cyclopropylmethyl, allyl and vinyl; wherein R^{12} and R^{13} is independently selected from methyl, ethyl and isopropyl; or a pharmaceutically-acceptable salt thereof.

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13. The composition of Claim 12 wherein said renin inhibitor compound is selected from compounds, their tautomers and pharmaceutically-acceptable salts thereof, of the group consisting of:

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15. A therapeutic method for treating hypertension or glaucoma, said method comprising administering to a hypertensive patient a therapeutically-effective amount of a compound of Formula I:

$$R^{13}$$
 N
 $(CH_2)_p$
 $S[O]_r$
 R^2
 R^3
 R^4
 R^6
 N
 R^6
 R^6
 R^8
 R^8

wherein each of R^1 and R^{11} is a group independently selected from hydrido, alkyl, alkylaminoalkyl and phenyl;

wherein p is a number selected from zero through five, inclusive; wherein r is a number selected from zero, one and two; wherein \mathbb{R}^2 is selected from hydrido and alkyl; wherein \mathbb{R}^3 is a group selected from hydrido,

cycloalkylalkyl, aralkyl and haloaralkyl; wherein each of ${\bf R}^4$ and ${\bf R}^6$ is a group independently selected from hydrido and methyl; wherein ${\bf R}^5$ is selected from

$$-(CH_2) \frac{\begin{bmatrix} R^9 \\ C \end{bmatrix}}{\begin{bmatrix} R^{10} \end{bmatrix}} - C \equiv C-V$$

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wherein V is selected from hydrido, alkyl, cycloalkyl, aryl and aralkyl; wherein each of R^9 and R^{10} is a group independently selected from hydrido, alkyl, alkenyl, alkynyl, cycloalkyl and aryl; wherein m is a number selected from zero through three; wherein n is a number selected from zero through three; wherein R^7 is a group selected from alkyl, cycloalkylalkyl and aralkyl; wherein R⁸ is a group selected from hydrido, alkyl, hydroxyalkyl, cycloalkyl, cycloalkylalkyl, alkenyl and haloalkenyl; wherein each of R^{12} and R^{13} is a group independently selected from hydrido, alkyl, cycloalkyl, cycloalkylalkyl, alkylacyl, aryl, aralkyl, haloaryl and haloaralkyl; and wherein any one of said R^1 through R^{13} groups having a substitutable position may be substituted with one or more groups selected from alkyl, hydroxy, hydroxyalkyl, halo, alkoxy, alkoxyalkyl and alkenyl; or a pharmaceutically-acceptable salt thereof.

16. The method of Claim 15 wherein each of R¹ and R¹¹ is independently selected from hydrido, methyl, ethyl, n-propyl, isopropyl, n-butyl, sec-butyl, isobutyl, tert-butyl, N,N'-dimethylaminomethyl, N,N'-diethylaminoethyl and phenyl;

wherein p is a number selected from zero through four, inclusive; wherein r is a number selected from zero, one and two; wherein R² is selected from hydrido and alkyl; wherein R³ is selected from hydrido, cycloalkylalkyl, phenylalkyl, halophenylalkyl, naphthylalkyl and halonaphthylalkyl; wherein each of R⁴ and R⁶ is independently selected from hydrido and methyl; wherein R⁵ is selected from

$$-(CH2) \frac{\begin{bmatrix} R^9 \\ C \end{bmatrix}}{\begin{bmatrix} C \end{bmatrix}} - C \equiv C-V$$

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wherein V is selected from hydrido, alkyl, phenyl and benzyl; wherein each of R^9 and R^{10} is independently selected from hydrido, alkyl, alkenyl, alkynyl, cycloalkyl and aryl; wherein m is a number selected from 15 zero through three; wherein n is a number selected from zero through three; wherein R^7 is selected from cyclohexylmethyl and benzyl, either one of which may be substituted with one or more groups selected from alkyl, hydroxy and alkoxy; wherein R⁸ is selected from hydrido, 20 alkyl, cycloalkyl, cycloalkylalkyl, hydroxyalkyl, alkenyl and haloalkenyl; and wherein each of R12 and R13 is independently selected from hydrido, alkyl, cycloalkyl, cycloalkylalkyl, alkanoyl, halophenyl, phenylalkyl, halophenylalkyl, naphthyl, halonaphthyl, naphthylalkyl 25 and halonaphthylalkyl; or a pharmaceutically-acceptable salt thereof.

17. The method of Claim 16 wherein each of R¹ and R¹¹ is independently selected from hydrido, methyl, ethyl, n-propyl and isopropyl; wherein p is a number selected from zero through three, inclusive; wherein r is

a number selected from zero, one and two; wherein R^2 is selected from hydrido, methyl, ethyl and n-propyl; wherein R^3 is selected from hydrido, cyclohexylmethyl, benzyl, phenylethyl, fluorobenzyl, fluorophenylethyl, chlorobenzyl, chlorophenylethyl, naphthylmethyl, naphthylethyl, fluoronaphthylmethyl and chloronaphthylmethyl; wherein each of R^4 and R^6 is independently selected from hydrido and methyl; wherein R^5 is selected from

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wherein V is selected from hydrido, alkyl, cycloalkyl, aryl and aralkyl; wherein m is a number selected from one 15 through three; wherein R⁷ is cyclohexylmethyl; wherein R⁸ is selected from methyl, ethyl, n-propyl, isopropyl, nbutyl, isobutyl, sec-butyl, tert-butyl, cyclopropyl, cyclobutyl, cyclopropylmethyl, cyclobutylmethyl, cyclohexylmethyl, allyl and vinyl; and wherein each of R12 20 and R13 is independently selected from hydrido, methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, sec-butyl, tert-butyl, cyclopropyl, cyclopropylmethyl, cyclopropylethyl, propylcarbonyl, ethylcarbonyl, methylcarbonyl, phenyl, benzyl, phenylethyl, 25 monochlorophenyl, dichlorophenyl, monofluorophenyl, difluorophenyl, monochlorophenylmethyl, monochlorophenylethyl, dichlorophenylmethyl, dichlorophenylethyl, naphthyl, monofluoronaphthyl, monochloronaphthyl, naphthylmethyl, naphthylethyl, 30 fluoronapthylmethyl and chloronaphthylethyl; or a pharmaceutically-acceptable salt thereof.

and R¹¹ is independently hydrido or methyl; wherein p is a number selected from zero through three, inclusive; wherein r is zero or two; wherein R² is selected from hydrido, methyl, ethyl and n-propyl; wherein R³ is selected from hydrido, cyclohexylmethyl, benzyl, phenylethyl, phenylpropyl, fluorobenzyl, fluorophenylethyl, chlorobenzyl, chlorophenylethyl, naphthylmethyl, fluoronaphthylmethyl and chloronaphthylmethyl; wherein each of R⁴ and R⁶ is hydrido; wherein R⁵ is selected from

$$--(CH_2)_m C \equiv C-V$$

wherein V is selected from hydrido and methyl; wherein m is one or two; wherein R⁷ is cyclohexylmethyl; wherein R⁸ is selected from ethyl, n-propyl, n-butyl, isobutyl, cyclopropyl, cyclobutyl, cyclopropylmethyl, allyl and vinyl; wherein each of R¹² and R¹³ is independently selected from hydrido, methyl, ethyl, n-propyl, isopropyl, cyclopropylmethyl, phenyl, benzyl, monochlorophenyl and dichlorophenyl; or a pharmaceutically-acceptable salt thereof.

19. The method of Claim 18 wherein said compound is of Formula II

$$R^{13} = \begin{bmatrix} R^{13} & 0 & R^{5} & H & OH \\ N & R^{12} & 0 & R^{7} & OH \end{bmatrix}$$

$$R^{13} = \begin{bmatrix} R^{13} & R^{13} & R^{13} & R^{12} & R^{13} & R^{12} & R^{12}$$

wherein r is zero or two; wherein q is two or three; wherein R² is selected from hydrido, methyl, ethyl and phenyl; wherein R³ is selected from hydrido, cyclohexylmethyl, benzyl, fluorobenzyl, chlorobenzyl,

fluoronaphthylmethyl and chloronaphthylmethyl; wherein each of R^4 and R^6 is hydrido; wherein R^5 is selected from

$$--(CH_2)_m C \equiv C-V$$

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wherein V is selected from hydrido and methyl; wherein m is one or two; wherein R⁷ is cyclohexylmethyl; wherein R⁸ is selected from n-propyl, isobutyl, cyclopropyl, cyclopropylmethyl, allyl and vinyl; wherein R¹² and R¹³ is independently selected from methyl, ethyl and isopropyl; or a pharmaceutically-acceptable salt thereof.

20. The method of Claim 18 wherein said compound is selected from compounds, their tautomers and pharmaceutically-acceptable salts thereof, of the group consisting of:

- 21. The method of Claim 20 wherein said compound is N-[1R*-[[1S,1R*-(cyclohexylmethyl)-2S*,3R*-dihydroxy-5-methylhexyl]amino]carbonyl]-3-butynyl]- α R*-[[[2-(dimethylamino)ethyl]sulfonyl] methyl]benzenepropanamide.
- 22. The method of Claim 15 for treating hypertension.
- 10 23. The method of Claim 15 for treating glaucoma.